D06

AMEMDMENTS TO THE CLAIMS

Please amend the claims to read as follows:

1. (Currently Amended) A method of inducing milk production in a mammal, the method comprising:

administering to the mammal, to induce lactation, a milk-secretion stimulating amount of:

- i) an estrogen-like agent (ELA); and
- [ii)]a progestational agent (PGA); wherein the first day of administration is defined as day 0; and
- ii) a {somatotropin} biologically active somatotropin (ST) {in said mammal,}; wherein the somatotropin is administered to the mammal to provide bioavailability of a milk-secretion stimulating amount beginning on day 0 and continuing for 20 days or more;

wherein the [first day of treatment is designated day 0 (zero)] mammal is a bovine selected from the group consisting of dairy heifers and dairy reproductive culls.

- (Currently Amended) The method of <u>claim</u> 1 wherein the ELA and PGA are administered as either a single sustained-release dose or are administered for approximately 5-12 days, beginning on day <u>0</u> [zero, and wherein the somatotropin is administered for at least 20 days from day 0].
- 3. (Currently Amended) The method of <u>claim</u> 2 wherein the ELA is administered at a dose of approximately 0.001 to 0.1 mg/kg/day and the PGA is administered at a dose of approximately .0025 to 0.25 mg/kg/day and is administered for approximately seven days.
- 4. (Currently Amended) The method of <u>claim</u> 2 wherein the somatotropin is administered for at least 30 days from day 0.
- 5. (Currently Amended) The method of claim 4 wherein the somatotropin is administered [on as] in approximately 4 doses.

D07

- 6. (Currently Amended) The method of <u>claim</u> 4 wherein the somatotropin is further administered in a sustained-release dose approximately every 14 days throughout lactation.
- 7. (Original) The method of claim 1, further comprising administering a milk-secretion enhancing amount of a glucocorticoid.
- 8. (Original) The method of claim 7 wherein the glucocorticoid is administered either on approximately day 10 to day 17 of the treatment or approximately 6 days after the final ELA administration.
- 9. (Original) The method of claim 7 wherein the glucocorticoid is dexamethasone administered at a dose of approximately 0.005 to 0.5 mg/kg on approximately day 10 to day 17 of the treatment.
- 10. (Original) The method of claim 1, further comprising subjecting the mammal to milk-stimulating photoperiods starting on, or before, day 0.
- 11. (Original) The method of claim 10 wherein said photoperiods:
 - a) comprise 12 consecutive hours of light and 12 consecutive hours of dark during each 24 hour period;
 - b) comprise progressively longer periods of light during each successive 24 hour period; or
 - c) comprise progressively shorter periods of light during each successive 24 hour period.

12.-13. (Cancelled)

- 14. (Currently Amended) The method of claim 1 further comprising providing physical stimulation of the [mammals] mammal's mammary gland region at least once daily for at least 7 consecutive days, starting on approximately day 7 of the treatment.
- 15. (Original) The method of claim 14 wherein the mammals mammary gland region is stimulated at least three times daily.

16. (Cancelled)

17. (Currently Amended) A method of inducing milk production in a mammal, the method comprising:

administering to the mammal, to induce lactation:

- i) an estrogen-like agent (ELA), subcutaneously, at a dose of approximately 0.007 to 0.7 mg/kg/body weight;
- ii) a progestational agent (PGA), subcutaneously, at a dose of approximately 0.0175 to 1.75 mg/kg/body weight/;
- iii) {glucocoriticoid} a glucocorticoid, intramuscularly, at a dose of approximately .005-.50 mg/kg/body weight; and
- iv) a [sematotropin] biologically active somatotropin [in the mammal], subcutaneously, at a dose of approximately 250-750 mg;

wherein ELA and PGA doses are each administered beginning on day 0; wherein the dexamethasone dose is administered on approximately day 10 to 17; wherein the first day of treatment is designated day 0 (zero); wherein the somatotropin is administered beginning on day 0 and periodically thereafter as required to maintain the bioavailability of a milk-secretion stimulating amount through at least day 20 of the treatment; and wherein the mammal is selected from the group consisting of dairy heifers and dairy reproductive culls.

18.-20. (Cancelled)

21. (Currently Amended) The method of claim {20} 1 wherein with the ELA is 17β-estradiol administered on day 0, as a slow-release pellet, at a dose of approximately 0.7 mg/kg body weight, wherein the PGA is natural progesterone administered on day 0, as a slow release pellet, at a dose of approximately 1.75 mg/kg body weight; wherein the somatotropin is administered as a sustained-release pellet approximately every two weeks throughout lactation; wherein the method further comprises administering, intramuscularly, dexamethasone on day 13, at a dose of approximately 0.05 mg/kg body weight; and wherein the treatment further comprises stimulating the mammary gland of the heifer or {cow} cull at least 2 times daily for at least seven consecutive days beginning on approximately day 7.

22.-25. (Cancelled)